Complete Summary

TITLE

Health plan members' experiences: percentage of parents of health plan members who reported their experiences with family centered care regarding their children with chronic conditions.

SOURCE(S)

CAHPS® Health Plan Survey and Reporting Kit 2002. Rockville (MD): Agency for Healthcare Research and Quality (AHRQ); 2002.

Welcome to the CAHPS Survey Users Network (CAHPS-SUN). [internet]. Rockville (MD): Agency for Healthcare Research and Quality (AHRQ); [cited 2004 Apr 15]. [3 p].

Brief Abstract

DESCRIPTION

This measure assesses the percentage of respondents who reported their experiences of family centered care for their children with chronic conditions. The "Family Center Care" composite measure is based on three composite measures ("Parents' experiences with the child's personal doctor or nurse," "Parents' experiences with shared decisionmaking," and "Parents' experiences with getting needed information about their child's care").

For children, the term "chronic conditions" refers to a wide range of physical, developmental, behavioral, and emotional health problems where the need for care is ongoing, and the services are of a type or at a level exceeding what children normally require.

RATIONALE

The Agency for Healthcare Research and Quality (AHRQ) (then called the Agency for Health Care Policy and Research, or AHCPR) initiated the CAHPS program in October 1995 to develop standardized survey tools for obtaining and reporting information on consumers' experiences with health care. The CAHPS consortium began by developing the CAHPS Health Plan Survey, an integrated set of carefully tested and standardized questionnaires and report formats that can be used to produce meaningful, reliable, and comparable information about the experiences of consumers enrolled in health plans.

The CAHPS Health Plan Survey is designed to generate information that consumers can use to choose health plans, that purchasers can use to assess the

value of services they buy, and that health plans can use to assess their performance and improve their products and services. As AHRQ had intended, the survey can be used with all types of health insurance consumers--including Medicaid recipients, Medicare beneficiaries, and those who are commercially insured--and across the full range of health care delivery systems, from fee-for-service to managed care plans. They instruments also capture information about special groups, including individuals with chronic conditions and disabilities and families with children.

The National Committee for Quality Assurance (NCQA) requires health plans to submit measures from the CAHPS Health Plan Survey as part of their HEDIS submission and for accreditation purposes.

Nationwide, the prevalence of children with chronic conditions is fairly low, with estimates ranging from 15-18 percent of all children under 18. However, some sponsors of CAHPS may serve a relatively higher proportion of children with chronic conditions than is found in the general population.

While their numbers may be small, financial impact, evidence of poor quality, equity, and the need for relevant information point to the need to measure and report on quality of care for children with chronic conditions.

PRIMARY CLINICAL COMPONENT

Health care; members' experiences; chronic conditions; family centered care (personal doctor or nurse, shared decisionmaking, getting needed information)

DENOMINATOR DESCRIPTION

Health plan members 17 years and younger, who have been enrolled in the health plan for 12 months or longer (commercial) or 6 months or longer (Medicaid), with no more than one break in enrollment of up to 45 days during the enrollment period, who have special health care needs (as determined by specific screening criteria included in the questionnaire), and whose parent answered the, "Family Centered Care," questions on the CAHPS 3.0 Health Plan Survey (Child Questionnaire)

NUMERATOR DESCRIPTION

The number of health plan members from the denominator whose parent reported their experiences with family centered care for their children with chronic conditions (see the related "Numerator Inclusions/Exclusions" field in the Complete Summary)

Evidence Supporting the Measure

PRIMARY MEASURE DOMAIN

Patient Experience

SECONDARY MEASURE DOMAIN

Access

EVIDENCE SUPPORTING THE MEASURE

A formal consensus procedure involving experts in relevant clinical, methodological, and organizational sciences

One or more research studies published in a National Library of Medicine (NLM) indexed, peer-reviewed journal

Evidence Supporting Need for the Measure

NEED FOR THE MEASURE

Overall poor quality for the performance measured

EVIDENCE SUPPORTING NEED FOR THE MEASURE

Bethell C, Lansky D, Hendryx M. The Robert Wood Johnson Foundation: national strategic indicators survey - summary report. Portland (OR): FACCT - The Foundation for Accountability; 2000 Sep 1. 82 p.

Bethell C. Measuring patient centered care across consumer relevant domains of quality: a report prepared for the Institute of Medicine Committee on the National Quality Report on Health Care. 2000 Jul 1.

State of Use of the Measure

STATE OF USE

Current routine use

CURRENT USE

Accreditation

Decision-making by businesses about health-plan purchasing
Decision-making by consumers about health plan/provider choice
External oversight/Department of Defense/TRICARE
External oversight/Medicaid
External oversight/State government program
Internal quality improvement
National health care quality reporting
Quality of care research

Application of Measure in its Current Use

CARE SETTING

Managed Care Plans

PROFESSIONALS RESPONSIBLE FOR HEALTH CARE

Nurses Physicians

LOWEST LEVEL OF HEALTH CARE DELIVERY ADDRESSED

Single Health Care Delivery Organizations

TARGET POPULATION AGE

Age less than or equal to 17 years

TARGET POPULATION GENDER

Either male or female

STRATIFICATION BY VULNERABLE POPULATIONS

Unspecified

Characteristics of the Primary Clinical Component

INCIDENCE/PREVALENCE

Nationwide, the prevalence of children with chronic conditions is fairly low, with estimates ranging from 15 to 18 percent of all children under 18.

EVIDENCE FOR INCIDENCE/PREVALENCE

CAHPS® Health Plan Survey and Reporting Kit 2002. Rockville (MD): Agency for Healthcare Research and Quality (AHRQ); 2002.

ASSOCIATION WITH VULNERABLE POPULATIONS

Children with special health care needs

EVIDENCE FOR ASSOCIATION WITH VULNERABLE POPULATIONS

CAHPS® Health Plan Survey and Reporting Kit 2002. Rockville (MD): Agency for Healthcare Research and Quality (AHRQ); 2002.

BURDEN OF ILLNESS

Unspecified

UTILIZATION

Unspecified

COSTS

Children with chronic conditions consume a preponderance of the health care dollars spent on children, with estimates ranging from 80 to 90 percent.

EVIDENCE FOR COSTS

CAHPS® Health Plan Survey and Reporting Kit 2002. Rockville (MD): Agency for Healthcare Research and Quality (AHRQ); 2002.

Institute of Medicine National Healthcare Quality Report Categories

IOM CARE NEED

End of Life Care Getting Better Living with Illness Staying Healthy

IOM DOMAIN

Patient-centeredness

Data Collection for the Measure

CASE FINDING

Users of care only

DESCRIPTION OF CASE FINDING

Health plan members 17 years or younger, who have been enrolled in the health plan for 12 months or longer (commercial) or 6 months or longer (Medicaid), with no more than one break in enrollment of up to 45 days during the enrollment period

DENOMINATOR SAMPLING FRAME

Enrollees or beneficiaries

DENOMINATOR (INDEX) EVENT

Patient Characteristic

DENOMINATOR INCLUSIONS/EXCLUSIONS

Inclusions

Health plan members 17 years and younger, who have been enrolled in the health plan for 12 months or longer (commercial) or 6 months or longer (Medicaid), with no more than one break in enrollment of up to 45 days during the enrollment period, who have special health care needs (as determined by specific screening criteria included in the questionnaire), and whose parent answered the, "Family Centered Care," questions on the CAHPS 3.0 Health Plan Survey (Child Questionnaire). Include refusals, non-response, and bad addresses/phone numbers.

Exclusions

- Individuals with coverage other than primary health coverage, such as dentalonly plan
- Deceased
- Ineligible (not enrolled in the plan)

NUMERATOR INCLUSIONS/EXCLUSIONS

Inclusions

The number of health plan members from the denominator whose parent reported their experiences with family centered care* for their children with chronic conditions

Includes all completed questionnaires; a questionnaire is considered complete if responses are available for 10 or more of a selected list of key CAHPS Health Plan Survey items.

*Family Centered Care:

- Parents' experiences with their children's personal doctor or nurse (2 questions)
- Parents' experiences with shared decisionmaking (4 questions)
- Parents' experiences with getting needed information about child's care (3 questions)

Exclusions Unspecified

DENOMINATOR TIME WINDOW

Time window precedes index event

NUMERATOR TIME WINDOW

Fixed time period

DATA SOURCE

Administrative data and patient survey

LEVEL OF DETERMINATION OF QUALITY

Not Individual Case

PRE-EXISTING INSTRUMENT USED

Children with Special Health Care Needs Screener, developed by the Child and Adolescent Health Measurement Initiative (CAHMI).

Computation of the Measure

SCORING

Non-weighted Score/Composite/Scale

INTERPRETATION OF SCORE

Better quality is associated with a higher score

ALLOWANCE FOR PATIENT FACTORS

Analysis by subgroup (stratification on patient factors) Case-mix adjustment

DESCRIPTION OF ALLOWANCE FOR PATIENT FACTORS

CAHPS recommends adjusting the data for respondent age, education, and general health status.

If the sample size is sufficient, responses may be analyzed for specific subpopulations, such as respondents with chronic conditions.

STANDARD OF COMPARISON

External comparison at a point in time External comparison of time trends Internal time comparison

Evaluation of Measure Properties

EXTENT OF MEASURE TESTING

The CAHPS Health Plan Survey has probably been tested more completely than any previously used consumer survey.

There are two different and complementary approaches to assessing the reliability and validity of a questionnaire (1) cognitive testing, which bases its assessments on feedback from interviews with people who are asked to react to the survey questions, and (2) psychometric testing, which bases its assessments on the

analysis of data collected by using the questionnaire. Although many existing consumer questionnaires about health care have been tested primarily or exclusively using a psychometric approach, the CAHPS team views the combination of cognitive and psychometric approaches as essential to producing the best possible survey instruments. Consequently, both methods have been included in the development of the CAHPS survey.

The cognitive testing method provided useful information on respondents 'perceptions of the response task, how respondents recalled and reported events, and how they interpreted specified reference periods. It also helped identify words that could be used to describe health care providers accurately and consistently across a range of consumers (e.g., commercially insured, Medicaid, fee-for-service, managed care, lower socioeconomic status [SES], middle SES, low literacy, higher literacy) and helped explore whether key words and concepts included in the core questions worked equally well in both English and Spanish.

The CAHPS consortium also tested each CAHPS reporting composite in focus groups with plan members. Cognitive interviews with consumers were conducted to ensure that the reporting composites and their labels were easily understood. Psychometric analyses using data collected during pilot tests were also conducted. These analyses indicated that both the composites and the items in each composite were reliable and valid measures of members' experiences. In addition, items in each reporting composite were tested and found to be internally consistent. For example, reliability coefficients (Cronbach's alpha) in one pilot test involving four health plans using the instrument that most resembled the final CAHPS 2.0 instrument ranged from a low of 0.68 for the "Getting Needed Care" composite to a high of 0.90 for the "How Well Doctors Communicate" composite. These composites are positively associated with members' ratings of overall care provided by doctors and nurses and ratings of health plans.

In addition, the CAHPS development team, together with researchers from the National Committee on Quality Assurance (NCQA), conducted a detailed comparative analysis of the items in the CAHPS questionnaire and NCQA's Member Satisfaction Survey (MSS) from the fall of 1997 to the spring of 1998. These questionnaires were merged to form the current CAHPS questionnaire. This testing is noteworthy because it was so extensive and because of the wide array of techniques used. These included focus groups, in-depth cognitive testing, pilot studies, methodological experiments, and large demonstration studies, such as the demonstrations in Washington State, Kansas, and New Jersey.

EVIDENCE FOR RELIABILITY/VALIDITY TESTING

Welcome to the CAHPS Survey Users Network (CAHPS-SUN). [internet]. Rockville (MD): Agency for Healthcare Research and Quality (AHRQ); [cited 2004 Apr 15]. [3 p].

Identifying Information

ORIGINAL TITLE

Family centered care.

MEASURE COLLECTION

CAHPS Health Plan Survey

MEASURE SET NAME

CAHPS 3.0 Health Plan Survey, Child Questionnaire

SUBMITTER

Agency for Healthcare Research and Quality

DEVELOPER

Agency for Healthcare Research and Quality CAHPS Consortium
Centers for Medicare and Medicaid Services

ADAPTATION

Measure was not adapted from another source.

RELEASE DATE

1997 Mar

REVISION DATE

2002 Oct

MEASURE STATUS

This is the current release of the measure.

SOURCE(S)

CAHPS® Health Plan Survey and Reporting Kit 2002. Rockville (MD): Agency for Healthcare Research and Quality (AHRQ); 2002.

Welcome to the CAHPS Survey Users Network (CAHPS-SUN). [internet]. Rockville (MD): Agency for Healthcare Research and Quality (AHRQ); [cited 2004 Apr 15]. [3 p].

MEASURE AVAILABILITY

The individual measure, "Family Centered Care," is published in the "CAHPS Health Plan Survey and Reporting Kit 2002." This Kit may be downloaded at the CAHPS-SUN Web site.

COMPANION DOCUMENTS

The following are available:

- Welcome to the CAHPS Survey Users Network (SUN). [Web site]. Available at www.cahps-sun.org.
- National CAHPS Benchmarking Database (NCBD). [Web site]. Rockville (MD): Agency for Healthcare Research and Quality (AHRQ); [cited 2003 Aug 21]. Available at http://ncbd.cahps.org/ or from the CAHPS-SUN Web site.
- What consumers say about the quality of their health plans and medical care.
 National CAHPS Benchmarking Database 2003 chartbook. Rockville (MD):
 Agency for Healthcare Research and Quality (AHRQ); 2003 Sep 1. 25 p.
 Available in Portable Document Format (PDF) from the NCBD Web site.
- Compare your health plan choices. Health plan quality from the consumer's point of view [adult guide]. Rockville (MD): CAHPS, Agency for Health Care Research and Quality; 17 p. Available from the CAHPS-SUN Web site.
- Compare your health plan choices. Health plan quality from the consumer's point of view [adult & child guide]. Rockville (MD): CAHPS, Agency for Healthcare Research and Quality; 25 p.Available from the <u>CAHPS-SUN</u> Web site.

NQMC STATUS

This NQMC summary was completed by ECRI on March 15, 2004. The information was verified by the measure developer on April 9, 2004.

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